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APPLICATION NO. FILING DATE FIRST NAMED INVENTOR ATTORNEY DOCKET NO. CONFIRMATION NO. 10/056,874 01/24/2002 Virginia W. Cornish 63711-A/JPW/GJG 3162 John P. White Cooper & Dunham LLP EPPERSON, JON D 1185 Avenue of the Americas New York, NY 10036 ART UNIT PAPER NUMBER 1639						
7590 06/16/2004 EXAMINER John P. White EPPERSON, JON D Cooper & Dunham LLP 1185 Avenue of the Americas ART UNIT PAPER NUMBER	APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
John P. White Cooper & Dunham LLP 1185 Avenue of the Americas ART UNIT PAPER NUMBER	10/056,874	01/24/2002	Virginia W. Cornish	63711-A/JPW/GJG	3162 ·	
Cooper & Dunham LLP 1185 Avenue of the Americas ART UNIT PAPER NUMBER	7	590 06/16/2004		EXAM	INER	
1185 Avenue of the Americas ART UNIT PAPER NUMBER New York NYC 10026	John P. White			EPPERSO	EPPERSON, JON D	
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New York, NY 10036				ART UNIT	PAPER NUMBER	
	New York, NY 10036			1639		

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)				
057 4-45 0	10/056,874	CORNISH, VIRGINIA W.				
Office Action Summary	Examiner	Art Unit				
	Jon D Epperson	1639				
The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply						
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 1 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely. - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication. - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).						
Status						
1) Responsive to communication(s) filed on	1) Responsive to communication(s) filed on					
2a) ☐ This action is FINAL . 2b) ☑ This	This action is FINAL. 2b)⊠ This action is non-final.					
3) Since this application is in condition for allowa	3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is					
closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.						
Disposition of Claims						
4)⊠ Claim(s) <u>1-54</u> is/are pending in the application.						
4a) Of the above claim(s) is/are withdrawn from consideration.						
5) Claim(s) is/are allowed.						
6) Claim(s) is/are rejected.) ☐ Claim(s) is/are rejected.					
7) Claim(s) is/are objected to.	Claim(s) is/are objected to.					
8) Claim(s) <u>1-54</u> are subject to restriction and/or election requirement.						
Application Papers						
9) The specification is objected to by the Examiner.						
10) ☐ The drawing(s) filed on is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.						
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).						
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).						
11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.						
Priority under 35 U.S.C. § 119						
12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).						
a) ☐ All b) ☐ Some * c) ☐ None of:						
1. Certified copies of the priority documents have been received.						
2. Certified copies of the priority documents have been received in Application No						
3. Copies of the certified copies of the priority documents have been received in this National Stage						
application from the International Bureau (PCT Rule 17.2(a)).						
* See the attached detailed Office action for a list of the certified copies not received.						
Attachment(s)						
1) Notice of References Cited (PTO-892)	4) 🔲 Interview Summary (
2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08) 4) Paper No(s)/Mail Date 5) Notice of Informal Patent Application (PTO-152)						
Paper No(s)/Mail Date 6) Other:						

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DETAILED ACTION

Election/Restriction

- 1. Restriction to one of the following inventions is required under 35 U.S.C. 121:
 - I. Claims 1-5 and 41-46, drawn to a product described as a "compound having the formula H1-Y-H2", classified variously in class 546, subclass 117 depending on the structure of the compound.
 - II. Claims 6-13, 47-50, drawn to a product described as "a complex between the compound of claim 1 and a fusion protein", classified variously in class 530, subclass 402+ depending on the structure of the complex.
 - III. Claims 14-22 and 51, drawn to a product described as a cell "comprising the complex of claim 6" or "comprising a DNA sequence which on transcription gives rise to a first fusion protein" classified variously in class 435, subclass 325+ depending on the cell type and the constituents therein.
 - IV. Claims 23-29, drawn to a method for "dimerizing two fusion proteins inside a cell" classified variously, for example, in class 435, subclass 7.2.
 - V. Claims 30-40 and 52-53, drawn to a method for "identifying a molecule that binds a known target in a cell from a pool of candidate molecules" classified variously, for example, in class 435, subclass 6.
 - VI. Claim 54, drawn to a product described as a "new protein cloned by the method of claim 53" classified variously, for example, in class 424, subclass 192.1.
- 2. The inventions are distinct, each from the other because of the following reasons:

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3. Groups I-VI represent separate and patentably distinct inventions. Groups IV-V are drawn to different methods and Groups I-III and VI are drawn to different products (i.e., e.g.,

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which are directed to different purposes, use different materials, recite different method or

process steps for the preparation of different product(s), screening of different characteristics,

such as different binding affinities, different biochemical reaction conditions, etc. or lead to

different final results). Therefore, the groups that describe these products and methods have

different issues regarding patentability and enablement, and represent patentably distinct subject

matter, which merits separate and burdensome searches. Art anticipating or rendering obvious

each of the above-identified groups respectively would not necessarily anticipate or render

obvious another group, because they are drawn to different inventions that have different

distinguishing features.

4. For Example, Groups IV-V represent patentably distinct methods. The methods are

distinct because they use different steps, require different reagents and/or will produce different

results. In this case, the method of Group V employs a library-screening step, which is not

required by the method of Group IV. As a result, Group V requires a different reagent (i.e., a

library) that is not required by Group IV. Therefore, Groups IV and V have different issues

regarding patentability and enablement and represent patentably distinct subject matter.

5. Likewise, Groups I-III and VI represent patentably distinct products. Groups I-III and VI

represent separate and patentably distinct products because they differ in respect to their

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properties, their use and the synthetic methodology for making them. For example, Group III is drawn to a "cell", which requires different reagents and/or materials than Groups I, II and VI (i.e., the other groups don't require cells). Likewise, Group II is drawn to a complex, which requires different reagents and/or materials than the groups I and VI (i.e., they don't require complexes). Finally Group VI is drawn to a "protein", which requires different reagents and/or materials than Group I. Therefore, art anticipating or rendering obvious each of the above-identified groups respectively would not necessarily anticipate or render obvious another group, because they are drawn to different inventions that have different distinguishing features and/or characteristics. Consequently, Groups I-III and VI have different issues regarding patentability

and enablement and represent patentably distinct subject matter.

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- In addition, if Applicants were to argue that any of Groups I-III and VI are somehow related to Groups IV-V as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP § 806.05(h)). In the instant case, the compounds can be used for a materially different process of using that product, such as in drug treatment therapies or in the materially different processes disclosed in Groups III and IV.
- 7. These inventions have acquired a separate status in the art as shown by their different classification and/or divergent subject matter. The different methods and products would require

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completely different searches in both the patent and non-patent databases, and there is no expectation that the searches would be coextensive. Therefore, this does create an undue search burden, and restriction for examination purposes as indicated is proper.

Species Election

- 8. This application contains claims directed to patentably distinct species of the claimed invention for Groups I-VI. Election is required as follows.
- 9. If applicant elects the invention of Group I, applicant is required to elect from the following patentably distinct species. Claim 1 is generic.

Subgroup 1: Species of compound having formula H1-Y-H2 (e.g., see claim 1)

Applicant must elect, for the purposes of search, a <u>single species</u> of compound having formula H1-Y-H2 wherein a <u>specific structure</u> is set forth, which clearly shows all of the atoms and bonds that are necessary to define the compound. Applicant should <u>not</u> use notations like X or R when identifying the elected structure because these letters represent groups with variable members and, as a result, more than one species would be erroneously elected (e.g., see compound in claim 5). Applicants must also indicate which parts of the molecule represent H1, Y and H2.

10. If applicant elects the invention of Group II, applicant is required to elect from the following patentably distinct species. Claim 6 is generic.

Subgroup 1: Species of compound having formula H1-Y-H2 (e.g., see claim 1)

Applicant must elect, for the purposes of search, a <u>single species</u> of compound having formula H1-Y-H2 wherein a <u>specific structure</u> is set forth, which clearly shows all of the atoms and bonds that are necessary to define the compound. Applicant should <u>not</u> use notations like X or R when identifying the elected structure because these letters represent groups with variable members and, as a result, more than one species would be

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erroneously elected (e.g., see compound in claim 5). Applicants must also indicate which parts of the molecule represent H1, Y and H2.

Subgroup 2: Species of fusion protein (e.g., see claim 6)

Applicant must elect, for the purposes of search, a <u>single species</u> of fusion protein e.g., eDHFR-LexA.

11. If applicant elects the invention of Group III, applicant is required to elect from the following patentably distinct species. Claim 14 is generic.

Subgroup 1: Species of compound having formula H1-Y-H2 if present (e.g., see claims 1, 6 and 14).

Applicant must elect, for the purposes of search, a <u>single species</u> of compound having formula H1-Y-H2 wherein a <u>specific structure</u> is set forth, which clearly shows all of the atoms and bonds that are necessary to define the compound. Applicant should <u>not</u> use notations like X or R when identifying the elected structure because these letters represent groups with variable members and, as a result, more than one species would be erroneously elected (e.g., see compound in claim 5). Applicants must also indicate which parts of the molecule represent H1, Y and H2.

Subgroup 2: Species of first fusion protein if present (e.g., see claim 15)

Applicant must elect, for the purposes of search, a <u>single species</u> of fusion protein e.g., eDHFR-LexA (see claim 20).

Subgroup 3: Species of second fusion protein if present (e.g., see claim 15)

Applicant must elect, for the purposes of search, a <u>single species</u> of fusion protein e.g., R61-B42 (see claim 20).

Subgroup 4: Species of cell (e.g., see claim 21)

Applicant must elect, for the purposes of search, a *single species* of cell e.g., yeast.

12. If applicant elects the invention of Group IV, applicant is required to elect from the following patentably distinct species. Claim 23 is generic.

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Subgroup 1: Species of compound having formula H1-Y-H2 (e.g., see claims 1, 23)

Applicant must elect, for the purposes of search, a <u>single species</u> of compound having formula H1-Y-H2 wherein a <u>specific structure</u> is set forth, which clearly shows all of the atoms and bonds that are necessary to define the compound. Applicant should <u>not</u> use notations like X or R when identifying the elected structure because these letters represent groups with variable members and, as a result, more than one species would be erroneously elected (e.g., see compound in claim 5). Applicants must also indicate which parts of the molecule represent H1, Y and H2.

Subgroup 2: Species of first fusion protein (e.g., see claim 29)

Applicant must elect, for the purposes of search, a <u>single species</u> of fusion protein e.g., eDHFR-LexA (see claim 20).

Subgroup 3: Species of second fusion protein (e.g., see claim 29)

Applicant must elect, for the purposes of search, a <u>single species</u> of fusion protein e.g., R61-B42 (see claim 20).

Subgroup 4: Species of cell (e.g., see claim 23)

Applicant must elect, for the purposes of search, a single species of cell e.g., yeast.

13. If applicant elects the invention of Group V, applicant is required to elect from the following patentably distinct species. Claim 30 is generic.

Subgroup 1: Species of covalent candidate (e.g., see claim 30)

Applicant must elect, for the purposes of search, a <u>single species</u> of covalent candidate. The election should result in a particularly defined core structure that is shared by all candidate molecules. In defining this core structure, all variable groups should be defined (i.e. all atoms and bonds shown) as much as possible. However, if no common core structure exists, a representative example must be elected.

Subgroup 2: Species of substrate (e.g., see claim 30)

Applicant must elect, for the purposes of search, a *single species* of substrate.

Subgroup 3: Species of target receptor (e.g., see claim 30)

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A. Known (e.g., see claim 30)

B. Unknown (e.g., see claim 38)

Applicant must elect, for the purposes of search, a <u>single species</u> of target receptor. Applicants must elect A or B above. In addition, Applicant must identify the target receptor if known or identify if it comes from a genomicDNA, cDNA or synthetic DNA if unknown (e.g., see claim 53).

Subgroup 4: Species of first fusion protein (e.g., see claim 30)

Applicant must elect, for the purposes of search, a <u>single species</u> of fusion protein e.g., eDHFR-LexA.

Subgroup 5: Species of second fusion protein (e.g., see claim 30)

Applicant must elect, for the purposes of search, a <u>single species</u> of fusion protein e.g., R61-B42.

Subgroup 6: Species of cell (e.g., see claim 31)

Applicant must elect, for the purposes of search, a single species of cell e.g., yeast.

Subgroup 7: Species of reporter gene (e.g., see claim 30)

Applicant must elect, for the purposes of search, a <u>single species</u> of reporter gene e.g., lacZ reporter gene.

Subgroup 7: Species of binding (e.g., see claim 30)

A. Competitive i.e., in presence of random small molecules for competitive binding (e.g., see claim 37)

B. Non-competitive i.e., <u>NOT</u> in presence of random small molecules for competitive binding (e.g., see claim 30).

Applicant must elect, for the purposes of search, a <u>single species</u> of binding. Applicants must elect A or B above.

14. If applicant elects the invention of Group VI, applicant is required to elect from the following patentably distinct species. Claim 54 is generic.

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Subgroup 1: Species of protein (e.g., see claim 54)

Applicant must elect, for the purposes of search, a <u>single species</u> of protein e.g., Applicants must provide the amino acid sequence.

- 15. <u>Please Note:</u> Applicants must disclose which claims read on the elected species (see paragraphs 19 and 20 below).
- 16. The species are distinct, each from the other, because their structures and modes of action are different. They would also differ in their reactivity and the starting materials from which they are made. For different species of method, the method steps for each species would differ. Moreover, the above species can be separately classified. Consequently, the species have different issues regarding patentability and represent patentably distinct subject matter. Therefore, this does create an undue search burden, and election for examination purposes as indicated is proper.
- 17. Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable.
- 18. Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance if the

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examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a a rejection under 35 U.S.C. 103(a) of the other invention.

- 19. Applicant is advised that a reply to this requirement <u>must include an identification of the</u>

 <u>species that is elected consonant with this requirement</u>, <u>and a listing of all claims readable</u>

 <u>thereon, including any claims subsequently added</u>. An argument that a claim is allowable or

 that all claims are generic is considered *nonresponsive* unless accompanied by an election.
- 20. Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, *applicant must indicate which are readable upon the elected species*. MPEP § 809.02(a).
- Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.43). Because the above restriction/election requirement is complex, a telephone call to applicants to request an oral election was not made. See MPEP § 812.01.
- 22. Applicant is reminded that upon cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the

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application. Any amendment of inventorship must be accompanied by a petition under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

- Applicant is also reminded that a 1 month (not less than 30 days) shortened statutory period will be set for response when a written requirement is made without an action on the merits. This period may be extended under the provisions of 37 CFR 1.136(a). Such action will not be an "action on the merits" for purposes of the second action final program, see MPEP 809.02(a).
- 24. Finally, Applicant is reminded that where applicant elects claims directed to a product, and a product claim is subsequently found allowable, withdrawn process claims that depend from or otherwise include all the limitations of the allowable product claim will be rejoined in accordance with the provisions of MPEP § 821.04. Process claims that depend from or otherwise include all the limitations of the patentable product will be entered as a matter of right if the amendment is presented prior to final rejection or allowance, whichever is earlier. Amendments submitted after final rejection are governed by 37 CFR 1.116; amendments submitted after allowance are governed by 37 CFR 1.312.

In the event of rejoinder, the requirement for restriction between the product claims and the rejoined process claims will be withdrawn, and the rejoined process claims will be fully examined for patentability in accordance with 37 CFR 1.104. Thus, to be allowable, the rejoined claims must meet all criteria for patentability including the requirements of 35 U.S.C. 101, 102, 103, and 112. Until an elected product claim is found allowable, an otherwise proper restriction

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requirement between product claims and process claims may be maintained. Withdrawn process claims that are not commensurate in scope with an allowed product claim will not be rejoined. See "Guidance on Treatment of Product and Process Claims in light of *In re Ochiai, In re Brouwer* and 35 U.S.C. § 103(b)," 1184 O.G. 86 (March 26, 1996). Additionally, in order to retain the right to rejoinder in accordance with the above policy, Applicant is advised that the process claims should be amended during prosecution either to maintain dependency on the product claims or to otherwise include the limitations of the product claims. **Failure to do so may result in a loss of the right to rejoinder.**

Further, note that the prohibition against double patenting rejections of 35 U.S.C. 121 does not apply where the restriction requirement is withdrawn by the examiner before the patent issues. See MPEP § 804.01.

Conclusion

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Jon D Epperson whose telephone number is (571) 272-0808. The examiner can normally be reached Monday-Friday from 9:00 to 5:30.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Andrew Wang can be reached on (571) 272-0811. The fax phone number for the organization where this application or proceeding is assigned is (703) 872-9306.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (571) 272-1600.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

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Jon D. Epperson, Ph.D. June 11, 2004

BENNETT CELSA PRIMARY EXAMINER

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